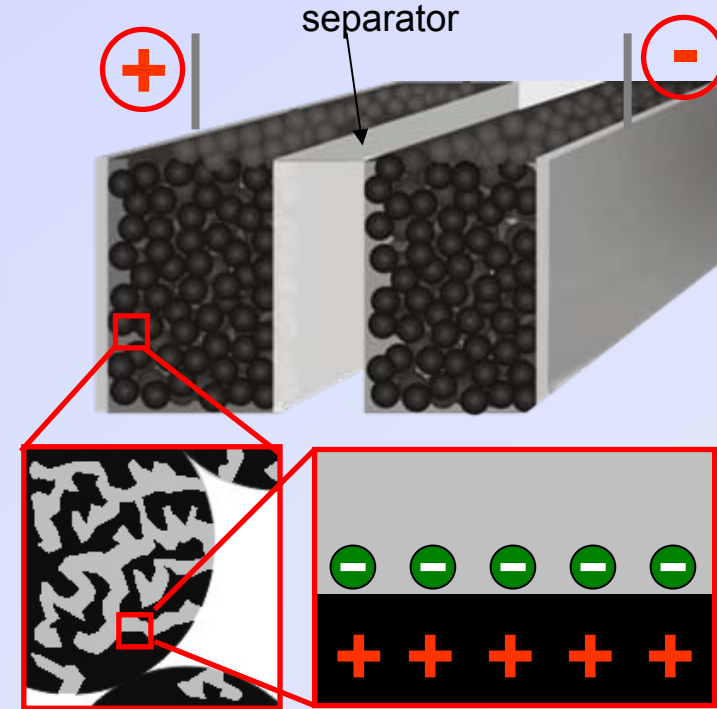
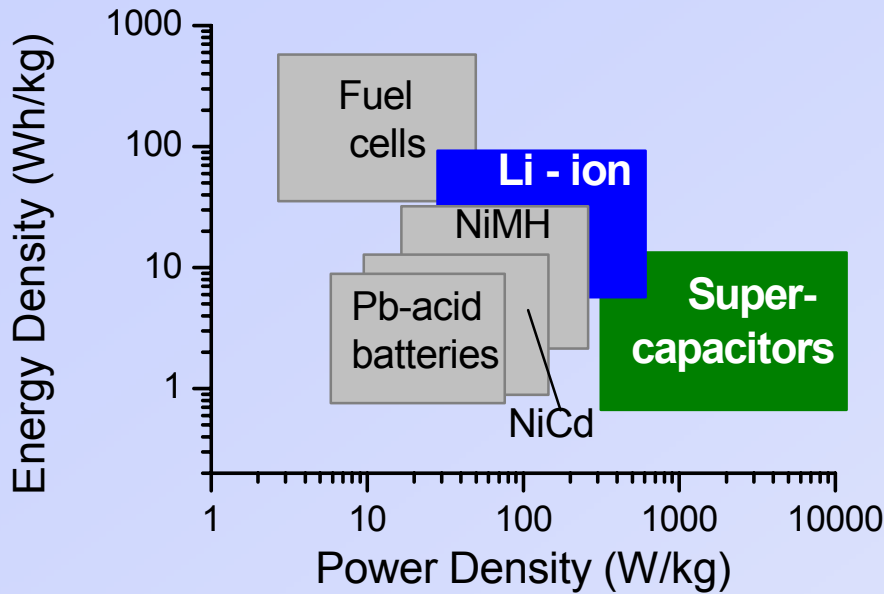


Integrated Supercapacitors For Nano-Morphic Systems

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Supercapacitors



- **Charge storage:**
 - electrical double-layer (EDLC)
 - fast and reversible faradaic redox reaction (pseudocapacitance)
- **Energy storage depends on the ability of electrode to adsorb electrolyte ions under the applied potential**

Supercapacitors

Advantages over batteries:

- **Higher power**
- **Faster charging** (seconds vs. hours)
- **Less harmful** (no heavy metals)
- **Offer long cycle life** (>1,000,000 cycles vs. 500)
- **High efficiency** (> 95%)
- **Easy to detect the state of charge**
- **No fundamental limit for the voltage** (voltage only restricted by the decomposition of the electrolyte)

Supercapacitors

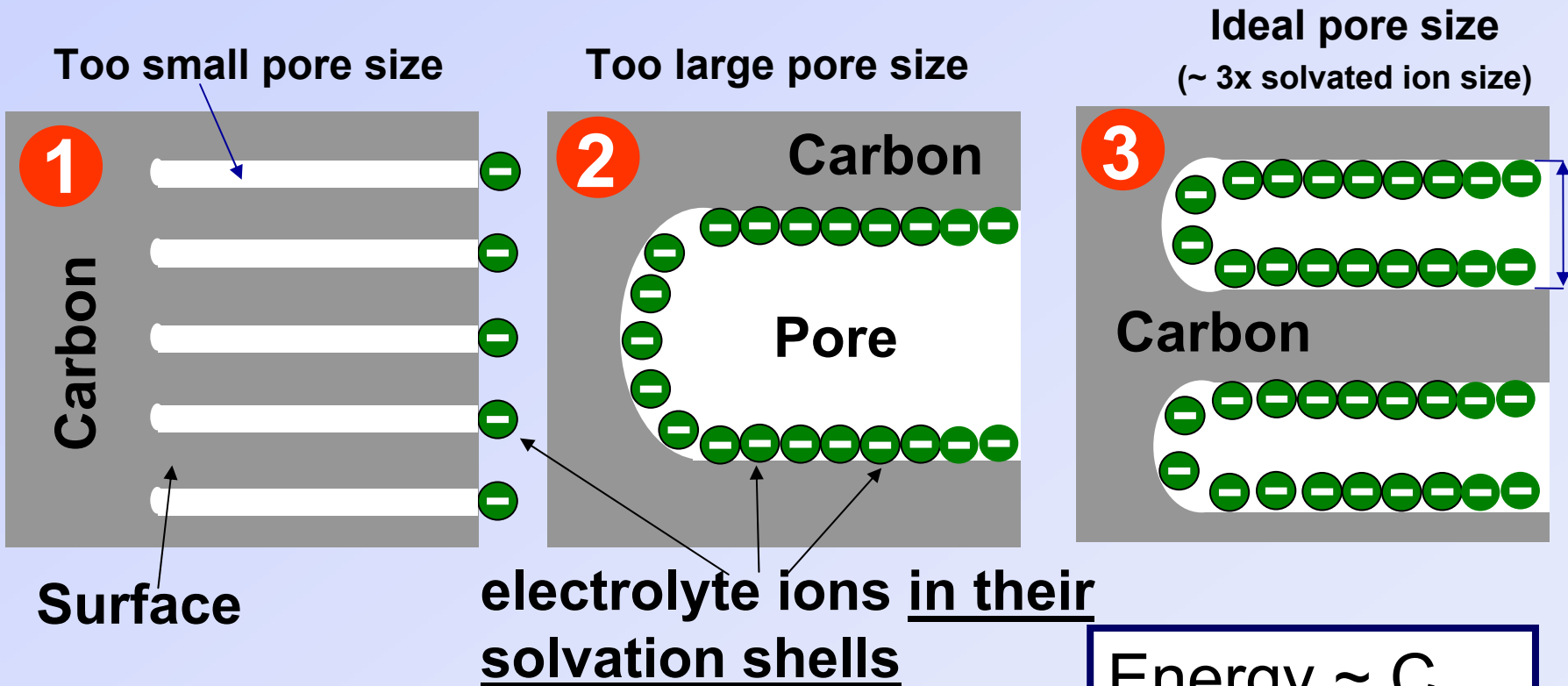
Why attractive for nano-morphic systems:

- **Provide peak power**
 - signal transmission
 - electroshock (e.g. destruction of cancer cells*; electro-therapy / cell stimulation)
 - activation (e.g. of bonding mechanism or
release of drugs that come near the target cell, etc.)
- **Opportunity to control ion concentration in the proximity**
 - sensing
 - curing cells (e.g. by controlling injection of specific ions)
 - selective ion adsorption depending on the ion size possible
- **Simple construction / easy miniaturization and integration**
- **Long life** (could be longer than human life)
- **In combination with fuel cells will provide the highest energy and power density**

Supercapacitors

How to increase the energy storage?

By better understanding the storage mechanisms!



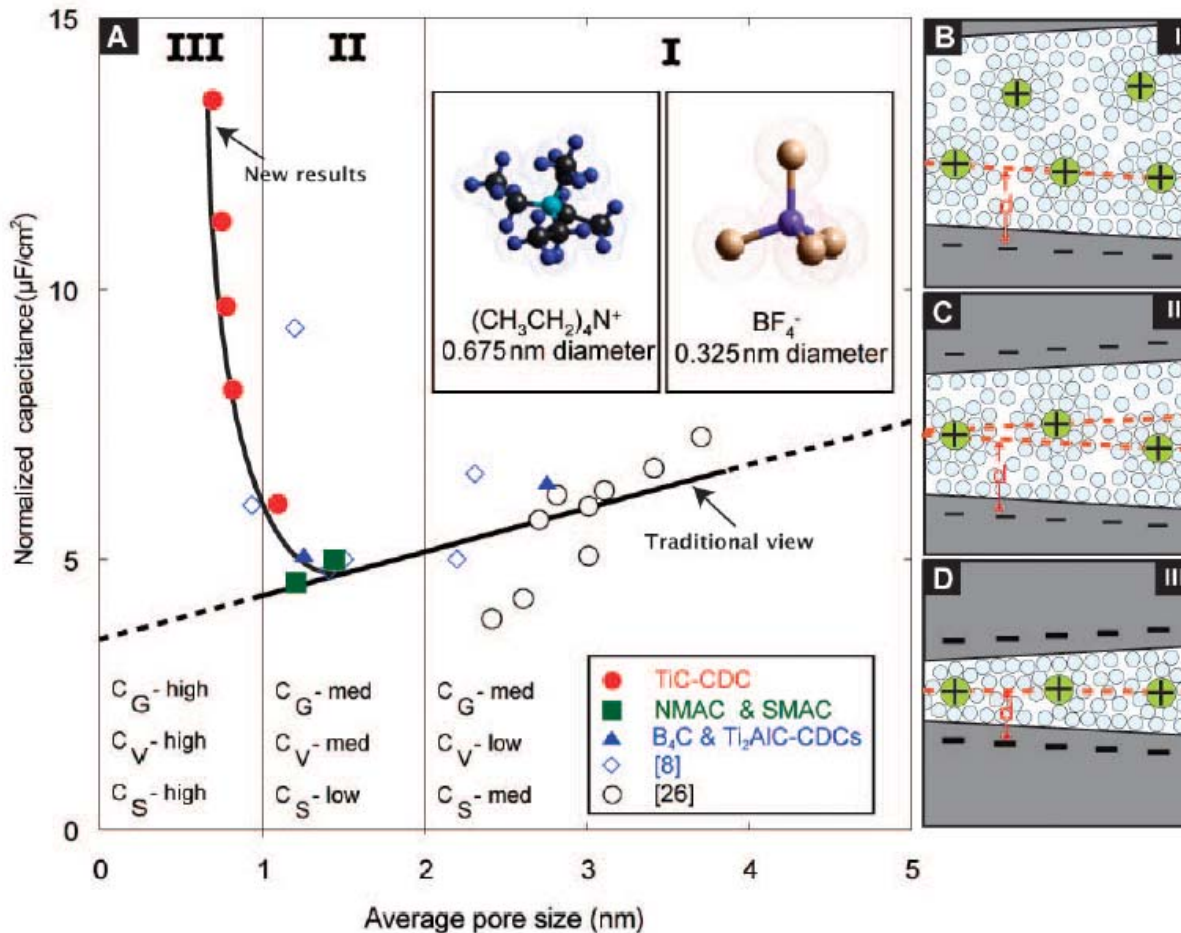
Traditional view

$$\text{Energy} \sim C$$

$$\text{Power} \sim \sqrt{\frac{C}{R}}$$

Supercapacitors

Energy storage in EDLC



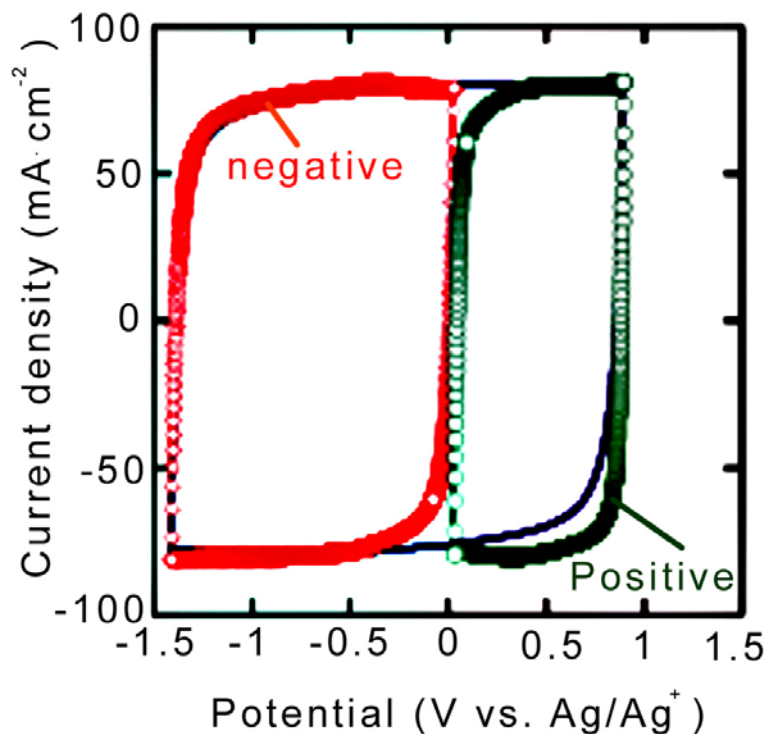
➤ Distortion of solvation shells in sub-nm pores results in enhanced ion storage capacities

➤ $C \sim 1/d$

Supercapacitors

How to achieve selective ion adsorption?

By better controlling the pore size and microstructure!



Cyclic Voltammogram taken at a scan rate of 20 mV/s in 3 electrode cell configuration

- Higher voltage drop on a negative electrode due to larger size of positive ions
- Difference in ions' size will affect their adsorption capacity if the pore size in electrode is tuned to dimensions of smaller ions

Active material: C synthesized from TiC and having the pore size of about 0.72 nm

The size of anion – about 0.48 nm
The size of cation – about 0.67 nm

Supercapacitors

What are the fundamental limits of energy storage ?

1. Empirical estimation:

Energy in supercapacitor device $E \approx \frac{CV^2}{2} \cdot \frac{1}{8}$ due to packaging, two C in a series etc.

Capacitance: (a) 6-30 uF/cm² in carbon; (b) up to 200 uF/cm² in functionalized carbon;
(c) up to 200 uF/cm² in transition metal oxides

If surface area = 2000 m²/g capacitance up to 4000 F/g could be reached

If max Voltage = 1V the Energy density $E = 70$ W·h/kg; if 3V, $E=630$ W·h/kg

2. Semi-empirical estimation:

Assume formation of a close-packed monolayer of solvated ions (1e; d = 1.5 nm): 0.4 C/m²

If could be achieved @ 1V: Capacitance = 4,000 uF/cm² and $E = 1,400$ W·h/kg

If could be achieved @ 10V: Capacitance = 400 uF/cm² and $E = 14,000$ W·h/kg

3. Fundamental limit:

If 1e/atom in electrode: eNV/2 per electrode (1/2 of batteries **but** V_{supercap} could be >2x higher)

Supercapacitors

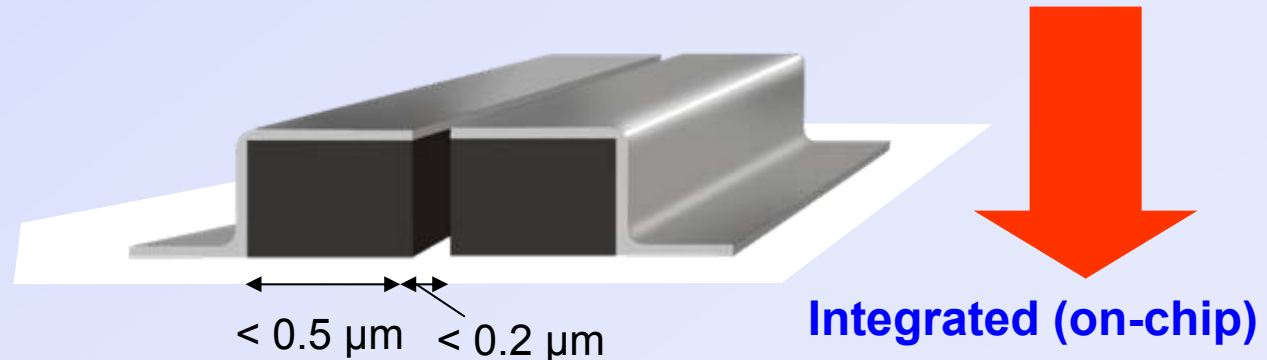
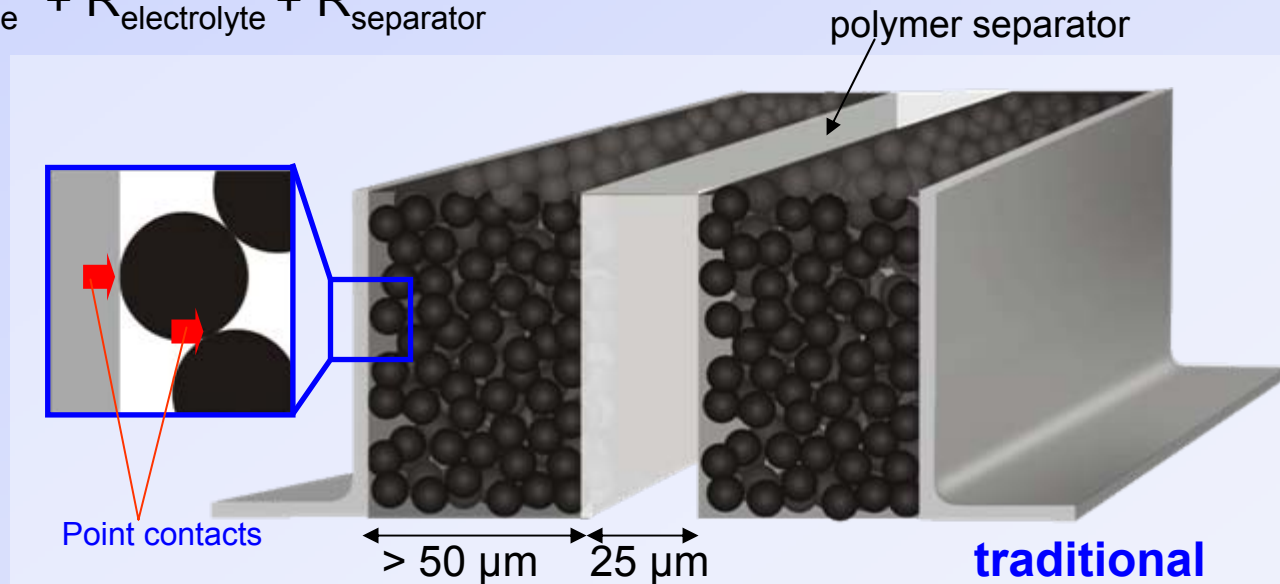
Will micro-fabrication affect supercapacitor performance ?

- Power achievable is related to the Equivalent Series Resistance (ESR)

$$ESR = R_{cc/electrode} + R_{electrode} + R_{electrolyte} + R_{separator}$$

By moving from traditional to integrated supercapacitor we:

- decrease $R_{cc/electrode}$ by > 100
- decrease $R_{electrode}$ by > 100
- decrease $R_{electrolyte}$ by > 100
- decrease $R_{separator}$ by > 100



Integrated supercap:

- $> 10x$ Power_(gravimetric)
- $> 20x$ Power_(volumetric)
- $< 10\%$ Charging time

Supercapacitors

- **Posses attractive properties** (higher power, faster charging, less harmful, very long cycle life, etc.)
- **High V might be possible** (voltage only restricted by the decomposition of the electrolyte)
- **Fundamentally do not have to exhibit low energy density** (state-of-the-art performance might be limited by our poor understanding of ion propagation and adsorption mechanisms and by materials synthesis technologies)
- **Going to micro- and nano- scale should increase power density by over 10 times**
- **Capable to control concentration of selected ions in the vicinity of the cells**
- **Might not need hermetic sealing** (can use electrolyte of the extracellular matrix)